Billing Code 4410-09-M

DEPARTMENT OF JUSTICE

DRUG ENFORCEMENT ADMINISTRATION

MANUFACTURER OF CONTROLLED SUBSTANCES

NOTICE OF REGISTRATION

JOHNSON MATTHEY PHARMA SERVICES

By Notice dated December 5, 2011, and published in the Federal Register on December 12, 2011, 76 FR 77257, Johnson Matthey Pharma Services, 70 Flagship Drive, North Andover, Massachusetts 01845, made application by renewal to the Drug Enforcement Administration (DEA) to be registered as a bulk manufacturer of the following basic classes of controlled substances:

Drug	Schedule
Amphetamine (1100)	II
Methylphenidate (1724)	II
Hydrocodone (9193)	II

The company plans to utilize this facility to manufacture small quantities of the listed controlled substances in bulk and to conduct analytical testing in support of the company's primary manufacturing facility in

West Deptford, New Jersey. The controlled substances

manufactured in bulk at this facility will be distributed

to the company's customers.

No comments or objections have been received. DEA has

considered the factors in 21 USC § 823(a) and determined

that the registration of Johnson Matthey Pharma Services to

manufacture the listed basic classes of controlled

substances is consistent with the public interest at this

time. DEA has investigated Johnson Matthey Pharma Services

to ensure that the company's registration is consistent

with the public interest. The investigation has included

inspection and testing of the company's physical security

systems, verification of the company's compliance with

state and local laws, and a review of the company's

background and history. Therefore, pursuant to 21 USC §

823(a), and in accordance with 21 CFR § 1301.33, the above

named company is granted registration as a bulk

manufacturer of the basic classes of controlled substances

listed.

> Joseph T. Rannazzisi
> Deputy Assistant Administrator
> Office of Diversion Control
> Drug Enforcement Administration

DATED: April 17, 2012